

? ds

Set	Items	Description
S1	9384424	S CANCER OR TUMOR OR TUMOUR OR MALIGNAN?
S2	3910277	S ANTIBOD?
S3	1034	S HM124 OR HM1(1N)24 OR BST2 OR BST(1N)2
S4	4985	S AU=KAWAI
S5	1179	S AU=MIHARA
S6	25	S AU=KOISHIHARA
S7	162	S S2(3N)S3
S8	119	RD (unique items)
S9	95	S S8 AND S1
S10	95	RD (unique items)
S11	4	S S4 AND S3
S12	1	S S5 AND S3
S13	15	S S6 AND S3
S14	17	S S11 OR S12 OR S13
S15	17	RD (unique items)
S16	1	S S15 NOT S10

? show files

[File 5] **Biosis Previews(R)** 1926-2007/Jul W1

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**File 5: BIOSIS has been enhanced with archival data. Please see HELP NEWS 5 for information.*

[File 34] **SciSearch(R) Cited Ref Sci** 1990-2007/Jul W2

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[File 35] **Dissertation Abs Online** 1861-2007/Jun

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[File 45] **EMCare** 2007/Jun W4

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[File 65] **Inside Conferences** 1993-2007/Jul 03

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[File 71] **ELSEVIER BIOBASE** 1994-2007/Jul W1

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2001 (c) Action Potential. All rights reserved.

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[File 144] **Pascal** 1973-2007/Jun W4

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[File 159] **Cancerlit** 1975-2002/Oct
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**File 159: Cancerlit is no longer updating. Please see HELP NEWS159.*

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[File 172] **EMBASE Alert** 2007/Jun 26
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[File 369] **New Scientist** 1994-2007/Jul W1
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[File 370] **Science** 1996-1999/Jul W3
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**File 370: This file is closed (no updates). Use File 47 for more current information.*

[File 399] **CA SEARCH(R)** 1967-2007/UD=14702
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[File 434] **SciSearch(R) Cited Ref Sci** 1974-1989/Dec
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[File 444] **New England Journal of Med.** 1985-2007/Jun W4
(c) 2007 Mass. Med. Soc. All rights reserved.

[File 467] **ExtraMED(tm)** 2000/Dec
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[File 654] **US PAT.FULL.** 1976-2007/JUN 28
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**File 654: IPCR/8 classification codes now searchable in 2006 records. For information about IC= index changes, see HELP NEWSIPCR.*

<!--StartFragment-->RESULT 11
ABB50295
ID ABB50295 standard; protein; 180 AA.
XX
AC ABB50295;
XX
DT 08-FEB-2002 (first entry)
XX
DE Bone marrow stromal antigen (BST-2) ovarian tumour marker protein, #80.
XX
KW Ovarian tumour marker gene; human; overexpression; upregulation;
KW epithelial tumour; cancer; diagnosis; prognosis; disease monitoring;
KW identification; serous cystadenoma; borderline serous tumour;
KW serous cystadenocarcinoma; mucinous cystadenocarcinoma;
KW mucinous cystadenoma; borderline mucinous tumour; endometrioid carcinoma;
KW undifferentiated carcinoma; clear cell adenocarcinoma; cystadenofibroma;
KW adenofibroma; Brenner tumour; serial analysis of gene expression; SAGE;
KW immune response pathway; cell proliferation regulation; protein folding;
KW membrane localised; secreted; therapeutic target; cytostatic;
KW gene therapy; vaccine.
XX
OS Homo sapiens.
XX
PN WO200175177-A2.
XX
PD 11-OCT-2001.
XX
PF 03-APR-2001; 2001WO-US010947.
XX
PR 03-APR-2000; 2000US-0194336P.
XX
PA (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Morin PJ, Sherman-Baust CA, Pizer ES, Hough CD;
XX
DR WPI; 2001-626450/72.
DR N-PSDB; ABA83121.
XX
PT Detecting and identifying ovarian tumor, identifying increased risk for
PT developing ovarian cancer, and determining effectiveness of ovarian
PT cancer treatment, by measuring expression level of ovarian tumor marker
PT gene.
XX
PS Claim 23; Page 124; 140pp; English.
XX
CC The invention relates to methods for diagnosing and prognosing ovarian
CC tumours in an individual via the detection and measurement of the
CC expression of ovarian tumour marker genes (ABA83081-ABA83122, ABA83180,
CC ABA83182 and ABA83184) or segments thereof (ABA83123-ABA83169, ABA83179,
CC ABA83181 and ABA83183). The methods of the invention are useful for
CC detecting an ovarian tumour in a patient, for identifying an individual
CC at increased risk for developing ovarian cancer, in prognostic tests for
CC assessing the relative severity of ovarian cancer, in tests for
CC monitoring a patient in remission from ovarian cancer and in tests for
CC monitoring disease status in a patient being treated for ovarian cancer.
CC The methods can additionally be used to identify a particular tumour as
CC being an ovarian tumour (i.e., an epithelial ovarian tumour selected from
CC serous cystadenoma, borderline serous tumour, serous cystadenocarcinoma,
CC mucinous cystadenoma, borderline mucinous tumour, mucinous
CC cystadenocarcinoma, endometrioid carcinoma, undifferentiated carcinoma,
CC clear cell adenocarcinoma, cystadenofibroma, adenofibroma and Brenner

CC tumour. The ovarian tumour marker genes of the invention were identified
CC using SAGE (serial analysis of gene expression) and were found to be
CC overexpressed in a broad variety of ovarian epithelial tumour cells
CC relative to normal ovarian epithelial cells. The marker genes are
CC implicated in immune response pathways, in the regulation of cell
CC proliferation and in protein folding, and many of these are membrane-
CC localised or secreted. In addition to their use as diagnostic and
CC prognostic markers, the ovarian tumour marker genes or their encoded
CC proteins may be used as therapeutic targets for the treatment and
CC prevention of ovarian cancer. Sequences ABB50257-ABB50299 represent
CC proteins encoded by ovarian tumour marker genes of the invention
XX

SQ Sequence 180 AA;

Query Match 100.0%; Score 889; DB 4; Length 180;
Best Local Similarity 100.0%; Pred. No. 2.2e-77;
Matches 180; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 MASTSYDYCRVPMEDGDKRCKLLLGIGILVLLIIVILGVPLIIFTIKANSEACRDGLRAV 60
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Db 1 MASTSYDYCRVPMEDGDKRCKLLLGIGILVLLIIVILGVPLIIFTIKANSEACRDGLRAV 60

Qy 61 MECRNVTHLLQQELTEAQKGFQDV EAQAATCNHTVMALMASLDAEKAQGQKKVEELEGEI 120
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Db 61 MECRNVTHLLQQELTEAQKGFQDV EAQAATCNHTVMALMASLDAEKAQGQKKVEELEGEI 120

Qy 121 TTLNHKLQDASAEVERLRENQVLSVRIADKKYYPSSQDSSAAAPQLLIVLLGLSALLQ 180
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Db 121 TTLNHKLQDASAEVERLRENQVLSVRIADKKYYPSSQDSSAAAPQLLIVLLGLSALLQ 180

<!--EndFragment-->